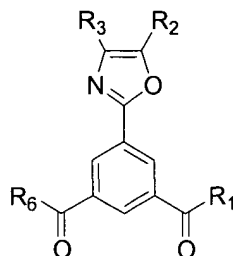


CLAIMS

What is claimed is:

1. A process for preparing a compound of the formula:



wherein:

R₁ is C₁-C₆ alkoxy or OH;

R₂ and R₃ are independently H, phenyl, or C₁-C₄ alkyl; or

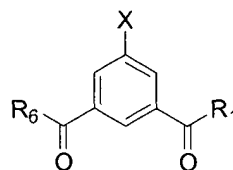
R₂ and R₃ and the carbons to which they are attached form a
benzo ring, which is optionally substituted with C₁-C₄ alkyl,
C₁-C₄ alkoxy, or dialkylamino; and

R₆ is C₁-C₆ alkoxy or NR₄R₅; wherein

R₄ and R₅ are independently C₁-C₆ alkyl;

comprising:

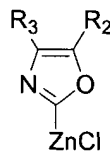
- forming a reaction mixture comprising a compound of
formula I:



I,

wherein X is Br, I, OTf, or OMs;

- a compound of formula II:



II,

a catalyst and at least one solvent.

2. A process according to claim 1, wherein the catalyst
is a transition metal catalyst.

3. A process according to claim 2 wherein the transition metal catalyst is $\text{Pd}(\text{PPh}_3)_4$, $\text{PdCl}_2(\text{PPh}_3)_2$, PdCl_2 , PdCl_2 and PPh_3 , or $\text{Pd}(\text{OCOCH}_3)_2$.

5

4. A process according to claim 3, wherein the catalyst is $\text{Pd}(\text{PPh}_3)_4$.

5. A process according to claim 1 wherein the method is conducted in the presence of at one additional polar, aprotic solvent.

6. A process according to claim 5, wherein the polar, aprotic solvent is tetrahydrofuran, tetramethyltetrahydrofuran, glyme, methyl t-butyl ether, or a mixture thereof.

7. A process according to claim 6, wherein the polar, aprotic solvent is tetrahydrofuran.

20

8. A process according to claim 1, wherein the reaction is performed at a temperature of from about 25°C to about the refluxing temperature of the solvent used.

9. A process according to claim 8 wherein the temperature is about 30°C to about 75°C .

10. A process according to claim 9, wherein the temperature is about 40°C to about 60°C .

30

11. A process according to claim 10, wherein the reaction mixture is formed by combining I, II and the catalyst, and any additional optional additives, at once or within a short time of each other.

35

12. A process according to claim 10, wherein the reaction mixture is formed over a period of about 0.5 hours to about 4 hours.

5 13. A process according to claim 12, wherein the time is about 1 hour to about 3 hours.

14. A process according to claim 13, wherein the time is about 1.5 hours to about 2.5 hours.

10

15. A process according to claim 1 wherein the transition metal catalyst is present in 0.01 to 20 mole percent, based on the amount of the compound of formula I.

15 16. A process according to claim 15, wherein the transition metal catalyst is present in 0.1 to 10 mole percent, based on the amount of the compound of formula I.

17. A process according to claim 16, wherein the
20 transition metal catalyst is present in 1 to 7 mole percent, based on the amount of the compound of formula I.

18. A process according to claim 17, wherein the reaction mixture is heated for about 24 hours.

25

19. A process according to claim 18, wherein the reaction mixture is heated for about 0.5 to about 8 hours.

20. A process according to claim 19, wherein the
30 reaction mixture is heated for about 0.5 to about 4 hours.

21. A process according to claim 20, wherein the reaction mixture is heated for about 0.5 to about 2.25 hours.

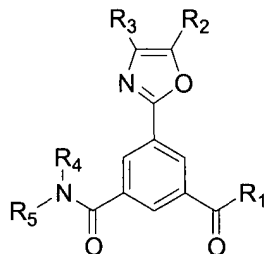
22. A process according to claim 1, wherein the compound of formula II is used in an excess from 1.001 to 10 equivalents, based on the compound of formula I.

5 23. A process according to claim 22, wherein the compound of formula II is used in an excess from 1.01 to 5 equivalents, based on the compound of formula I.

24. A process according to claim 23, wherein the
10 compound of formula II is used in an excess of 3 equivalents, based on the compound of formula I.

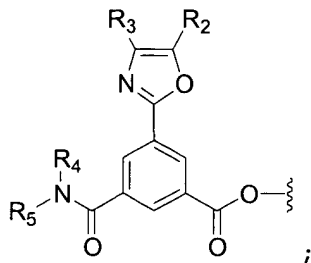
25. A process according to claim 1, wherein
X is Br;
15 R_2 and R_3 are independently H, methyl or ethyl;
 R_6 is NR_4R_5 ; wherein
 R_4 and R_5 are both C_3 alkyl; and
 R_1 is C_1 - C_4 alkyl.

20 26. A compound of the formula:



wherein:

R_1 is OH, imidazolyl, halogen, $-OC(O)CH_3$, $-OC(O)CF_3$;



25 R_2 and R_3 are independently H or C_1 - C_4 alkyl; and
 R_4 and R_5 are independently C_1 - C_6 alkyl.

27. A compound according to claim 26, wherein
R₂ and R₃ are independently H or methyl.

5 28. A compound according to claim 27, wherein R₄ and R₅
are both C₃ alkyl.

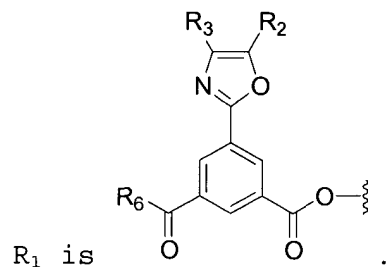
29. A compound according to claim 28, wherein
R₁ is OH.

10

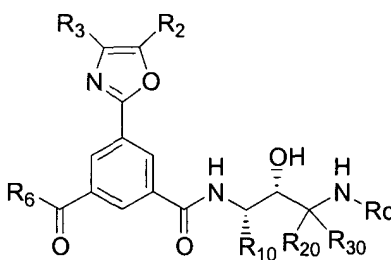
30. A compound according to claim 28, wherein
R₁ is C₁-C₄ alkoxy.

31. A compound according to claim 28, wherein
15 R₁ is chloro.

32. A compound according to claim 28, wherein



20 33. A process for preparing compounds of the formula:



wherein

R₁₀ is -(CH₂)₁₋₂-S(O)₀₋₂-(C₁-C₆ alkyl), or

25 C₁-C₁₀ alkyl optionally substituted with 1, 2, or 3 groups
independently selected from halogen, -OH, =O, -SH,
-C≡N, -CF₃, -C₁-C₃ alkoxy, amino, mono- or

dialkylamino, $-N(R)C(O)R'-$, $-OC(=O)-$ amino and $-OC(=O)-$ mono- or dialkylamino, or
 C_2-C_6 alkenyl or C_2-C_6 alkynyl, each of which is optionally substituted with 1, 2, or 3 groups independently
5 selected from halogen, $-OH$, $-SH$, $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy, amino, and mono- or dialkylamino, or
aryl, heteroaryl, heterocyclyl, $-C_1-C_6$ alkyl-aryl, $-C_1-C_6$ alkyl-heteroaryl, or $-C_1-C_6$ alkyl-heterocyclyl, where
the ring portions of each are optionally substituted
10 with 1, 2, 3, or 4 groups independently selected
from halogen, $-OH$, $-SH$, $-C\equiv N$, $-NR_{105}R'_{105}$, $-CO_2R$, $-N(R)COR'$, or $-N(R)SO_2R'$, $-C(=O)-(C_1-C_4)$ alkyl, $-SO_2-$ amino, $-SO_2-$ mono or dialkylamino, $-C(=O)-$ amino,
 $-C(=O)-$ mono or dialkylamino, $-SO_2-(C_1-C_4)$ alkyl, or
15 C_1-C_6 alkoxy optionally substituted with 1, 2, or 3
groups which are independently selected from
halogen, or
 C_3-C_7 cycloalkyl optionally substituted with 1, 2, or
3 groups independently selected from halogen,
20 $-OH$, $-SH$, $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy, amino, $-C_1-C_6$
alkyl and mono- or dialkylamino, or
 C_1-C_{10} alkyl optionally substituted with 1, 2, or 3
groups independently selected from halogen, $-OH$, $-SH$, $-C\equiv N$, $-CF_3$, $-C_1-C_3$ alkoxy, amino, mono-
25 or dialkylamino and $-C_1-C_3$ alkyl, or
 C_2-C_{10} alkenyl or C_2-C_{10} alkynyl each of which is
optionally substituted with 1, 2, or 3 groups
independently selected from halogen, $-OH$, $-SH$,
 $-C\equiv N$, $-CF_3$, C_1-C_3 alkoxy, amino, C_1-C_6 alkyl and
30 mono- or dialkylamino; and the heterocyclyl
group is optionally further substituted with
oxo;
R and R' independently are hydrogen, C_1-C_{10} alkyl, C_1-
 C_{10} alkylaryl or C_1-C_{10} alkylheteroaryl;

R₂₀ is selected from the group consisting of H; C₁-C₆ alkyl, optionally substituted with 1, 2, or 3 substituents that are independently selected from the group consisting of C₁-C₃ alkyl, halogen, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b}; -(CH₂)₀₋₄-aryl; -(CH₂)₀₋₄-heteroaryl; C₂-C₆ alkenyl; C₂-C₆ alkynyl; -CONR_{N-2}R_{N-3}; -SO₂NR_{N-2}R_{N-3}; -CO₂H; and -CO₂-(C₁-C₄ alkyl); wherein

R_{1-a} and R_{1-b} are independently -H or C₁-C₆ alkyl;

R₃₀ is selected from the group consisting of H; C₁-C₆ alkyl, optionally substituted with 1, 2, or 3 substituents independently selected from the group consisting of C₁-C₃ alkyl, halogen, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR_{1-a}R_{1-b}; -(CH₂)₀₋₄-aryl; -(CH₂)₀₋₄-heteroaryl; C₂-C₆ alkenyl; C₂-C₆ alkynyl; -CO-NR_{N-2}R_{N-3}; -SO₂-NR_{N-2}R_{N-3}; -CO₂H; and -CO-O-(C₁-C₄ alkyl);

or

R₂₀, R₃₀ and the carbon to which they are attached form a carbocycle of three thru seven carbon atoms, wherein one carbon atom is optionally replaced by a group selected from-O-, -S-, -SO₂-, or -NR_{N-2}-;

R_{N-2} and R_{N-3} at each occurrence are independently selected from the group consisting of -C₁-C₈ alkyl optionally substituted with 1, 2, or 3 groups independently selected from the group consisting of -OH, -NH₂, phenyl and halogen; -C₃-C₈ cycloalkyl; -(C₁-C₂ alkyl)-(C₃-C₈ cycloalkyl); -(C₁-C₆ alkyl)-O-(C₁-C₃ alkyl); -C₂-C₆ alkenyl; -C₂-C₆ alkynyl; -C₁-C₆ alkyl chain with one double bond and one triple bond; aryl; heteroaryl; heterocycloalkyl;

or

R_{N-2}, R_{N-3} and the nitrogen to which they are attached form a 5, 6, or 7 membered heterocycloalkyl or heteroaryl group, wherein said heterocycloalkyl or heteroaryl group is optionally fused to a benzene, pyridine, or pyrimidine ring, and said groups are unsubstituted or substituted with 1, 2, 3, 4, or 5 groups that at

each occurrence are independently C₁-C₆ alkyl, C₁-C₆ alkoxy, halogen, halo C₁-C₆ alkyl, halo C₁-C₆ alkoxy, -CN, -NO₂, -NH₂, NH(C₁-C₆ alkyl), N(C₁-C₆ alkyl)(C₁-C₆ alkyl), -OH, -C(O)NH₂, -C(O)NH(C₁-C₆ alkyl),
5 -C(O)N(C₁-C₆ alkyl)(C₁-C₆ alkyl), C₁-C₆ alkoxy C₁-C₆ alkyl, C₁-C₆ thioalkoxy, and C₁-C₆ thioalkoxy C₁-C₆ alkyl;
R_C is hydrogen, -(CR₂₄₅R₂₅₀)₀₋₄-aryl, -(CR₂₄₅R₂₅₀)₀₋₄-heteroaryl, -(CR₂₄₅R₂₅₀)₀₋₄-heterocyclyl, -(CR₂₄₅R₂₅₀)₀₋₄-aryl-heteroaryl, -(CR₂₄₅R₂₅₀)₀₋₄-aryl-heterocyclyl, -(CR₂₄₅R₂₅₀)₀₋₄-aryl-aryl,
10 -(CR₂₄₅R₂₅₀)₀₋₄-heteroaryl-aryl, -(CR₂₄₅R₂₅₀)₀₋₄-heteroaryl-heterocyclyl, -(CR₂₄₅R₂₅₀)₀₋₄-heteroaryl-heteroaryl, -(CR₂₄₅R₂₅₀)₀₋₄-heterocyclyl-heteroaryl, -(CR₂₄₅R₂₅₀)₀₋₄-heterocyclyl-heterocyclyl, -(CR₂₄₅R₂₅₀)₀₋₄-heterocyclyl-aryl,
15 -[C(R₂₅₅)(R₂₆₀)]₁₋₃-CO-N-(R₂₅₅)₂, -CH(aryl)₂, -CH(heteroaryl)₂, -CH(heterocyclyl)₂, -CH(aryl)(heteroaryl), -(CH₂)₀₋₁-CH((CH₂)₀₋₆-OH)-(CH₂)₀₋₁-aryl, -(CH₂)₀₋₁-CH((CH₂)₀₋₆-OH)-(CH₂)₀₋₁-heteroaryl, -CH(-aryl or -heteroaryl)-CO-O(C₁-C₄ alkyl), -CH(-CH₂-OH)-CH(OH)-phenyl-NO₂, (C₁-C₆ alkyl)-O-(C₁-C₆ alkyl)-OH; -CH₂-NH-CH₂-CH(-O-CH₂-CH₃)₂, -(CH₂)₀₋₆-C(=NR₂₃₅)(NR₂₃₅R₂₄₀), or
C₁-C₁₀ alkyl optionally substituted with 1, 2, or 3 groups independently selected from the group consisting of R₂₀₅, -OC=ONR₂₃₅R₂₄₀, -S(=O)₀₋₂(C₁-C₆ alkyl), -SH,
25 -NR₂₃₅C=ONR₂₃₅R₂₄₀, -C=ONR₂₃₅R₂₄₀, and -S(=O)₂NR₂₃₅R₂₄₀, or -(CH₂)₀₋₃-(C₃-C₈) cycloalkyl wherein the cycloalkyl is optionally substituted with 1, 2, or 3 groups independently selected from the group consisting of R₂₀₅, -CO₂H, and -CO₂-(C₁-C₄ alkyl), or
30 cyclopentyl, cyclohexyl, or cycloheptyl ring fused to aryl, heteroaryl, or heterocyclyl wherein one, two or three carbons of the cyclopentyl, cyclohexyl, or cycloheptyl is optionally replaced with a heteroatom independently selected from NH, NR₂₁₅, O, or S(=O)₀₋₂,
35 and wherein the cyclopentyl, cyclohexyl, or cycloheptyl group can be optionally substituted with

one or two groups that are independently R_{205} , $=O$,
-CO-NR₂₃₅R₂₄₀, or -SO₂-(C₁-C₄ alkyl), or
C₂-C₁₀ alkenyl or C₂-C₁₀ alkynyl, each of which is
optionally substituted with 1, 2, or 3 R_{205} groups,
5 wherein
each aryl and heteroaryl is optionally substituted with
1, 2, or 3 R_{200} , and wherein each heterocyclyl is
optionally substituted with 1, 2, 3, or 4 R_{210} ;
 R_{200} at each occurrence is independently selected from -OH,
10 -NO₂, halogen, -CO₂H, C≡N, -(CH₂)₀₋₄-CO-NR₂₂₀R₂₂₅, -(CH₂)₀₋₄-
CO-(C₁-C₁₂ alkyl), -(CH₂)₀₋₄-CO-(C₂-C₁₂ alkenyl), -(CH₂)₀₋₄-
CO-(C₂-C₁₂ alkynyl), -(CH₂)₀₋₄-CO-(C₃-C₇ cycloalkyl), -
(CH₂)₀₋₄-CO-aryl, -(CH₂)₀₋₄-CO-heteroaryl, -(CH₂)₀₋₄-CO-
heterocyclyl, -(CH₂)₀₋₄-CO-O-R₂₁₅, -(CH₂)₀₋₄-SO₂-NR₂₂₀R₂₂₅, -
15 (CH₂)₀₋₄-SO-(C₁-C₈ alkyl), -(CH₂)₀₋₄-SO₂-(C₁-C₁₂ alkyl), -
(CH₂)₀₋₄-SO₂-(C₃-C₇ cycloalkyl), -(CH₂)₀₋₄-N(H or R_{215})-CO-O-
R₂₁₅, -(CH₂)₀₋₄-N(H or R_{215})-CO-N(R_{215})₂, -(CH₂)₀₋₄-N-CS-
N(R_{215})₂, -(CH₂)₀₋₄-N(-H or R_{215})-CO-R₂₂₀, -(CH₂)₀₋₄-NR₂₂₀R₂₂₅,
-(CH₂)₀₋₄-O-CO-(C₁-C₆ alkyl), -(CH₂)₀₋₄-O-P(O)-(OR₂₄₀)₂,
20 -(CH₂)₀₋₄-O-CO-N(R_{215})₂, -(CH₂)₀₋₄-O-CS-N(R_{215})₂, -(CH₂)₀₋₄-O-
(R_{215}), -(CH₂)₀₋₄-O-(R_{215})-COOH, -(CH₂)₀₋₄-S-(R_{215}), -(CH₂)₀₋₄-
O-(C₁-C₆ alkyl optionally substituted with 1, 2, 3, or 5 -
F), C₃-C₇ cycloalkyl, -(CH₂)₀₋₄-N(H or R_{215})-SO₂-R₂₂₀, -(CH₂)₀₋₄-
C₃-C₇ cycloalkyl, or
25 C₁-C₁₀ alkyl optionally substituted with 1, 2, or 3 R_{205}
groups, or
C₂-C₁₀ alkenyl or C₂-C₁₀ alkynyl, each of which is
optionally substituted with 1 or 2 R_{205} groups,
wherein
30 the aryl and heteroaryl groups at each occurrence are
optionally substituted with 1, 2, or 3 groups that
are independently R_{205} , R_{210} , or
C₁-C₆ alkyl substituted with 1, 2, or 3 groups that
are independently R_{205} or R_{210} , and wherein

the heterocyclyl group at each occurrence is optionally substituted with 1, 2, or 3 groups that are independently R_{210} ;

R_{205} at each occurrence is independently selected from C_1 - C_6 alkyl, halogen, -OH, -O-phenyl, -SH, -C \equiv N, -CF $_3$, C_1 - C_6 alkoxy, NH $_2$, NH(C_1 - C_6 alkyl) or N-(C_1 - C_6 alkyl)(C_1 - C_6 alkyl);

R_{210} at each occurrence is independently selected from halogen, C_1 - C_6 alkoxy, C_1 - C_6 haloalkoxy, -NR $_{220}$ R $_{225}$, OH, C \equiv N, -CO-(C_1 - C_4 alkyl), -SO $_2$ -NR $_{235}$ R $_{240}$, -CO-NR $_{235}$ R $_{240}$, -SO $_2$ -(C_1 - C_4 alkyl), =O, or C_1 - C_6 alkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl or C_3 - C_7 cycloalkyl, each of which is optionally substituted with 1, 2, or 3 R_{205} groups;

R_{215} at each occurrence is independently selected from C_1 - C_6 alkyl, -(CH $_2$) $_{0-2}$ -(aryl), C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, C_3 - C_7 cycloalkyl, and -(CH $_2$) $_{0-2}$ -(heteroaryl), -(CH $_2$) $_{0-2}$ -(heterocyclyl), wherein the aryl group at each occurrence is optionally substituted with 1, 2, or 3 groups that are independently R_{205} or R_{210} , and wherein the heterocyclyl and heteroaryl groups at each occurrence are optionally substituted with 1, 2, or 3 R_{210} ;

R_{220} and R_{225} at each occurrence are independently selected from -H, - C_3 - C_7 cycloalkyl, -(C_1 - C_2 alkyl)-(C_3 - C_7 cycloalkyl), -(C_1 - C_6 alkyl)-O-(C_1 - C_3 alkyl), - C_2 - C_6 alkenyl, - C_2 - C_6 alkynyl, - C_1 - C_6 alkyl chain with one double bond and one triple bond, -aryl, -heteroaryl, and -heterocyclyl, or - C_1 - C_{10} alkyl optionally substituted with -OH, -NH $_2$ or halogen, wherein the aryl, heterocyclyl and heteroaryl groups at each occurrence are optionally substituted with 1, 2, or 3 R_{270} groups

R_{235} and R_{240} at each occurrence are independently H, or C_1 - C_6 alkyl;

R₂₄₅ and R₂₅₀ at each occurrence are independently selected from
-H, C₁-C₄ alkyl, C₁-C₄ alkylaryl, C₁-C₄ alkylheteroaryl, C₁-
C₄ hydroxyalkyl, C₁-C₄ alkoxy, C₁-C₄ haloalkoxy, -(CH₂)₀₋₄-
C₃-C₇ cycloalkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, and phenyl;
5 or

R₂₄₅ and R₂₅₀ are taken together with the carbon to which they
are attached to form a carbocycle of 3, 4, 5, 6, or 7
carbon atoms, where one carbon atom is optionally
replaced by a heteroatom selected from -O-, -S-, -SO₂-,
10 and -NR₂₂₀-;

R₂₅₅ and R₂₆₀ at each occurrence are independently selected from
-H, -(CH₂)₁₋₂-S(O)₀₋₂-(C₁-C₆ alkyl), -(C₁-C₄ alkyl)-aryl,
-(C₁-C₄ alkyl)-heteroaryl, -(C₁-C₄ alkyl)-heterocyclyl, -
aryl, -heteroaryl, -heterocyclyl, -(CH₂)₁₋₄-R₂₆₅-(CH₂)₀₋₄-
15 aryl, -(CH₂)₁₋₄-R₂₆₅-(CH₂)₀₋₄-heteroaryl, -(CH₂)₁₋₄-R₂₆₅-(CH₂)₀₋₄-
heterocyclyl, or

C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl or -(CH₂)₀₋₄-C₃-C₇
cycloalkyl, each of which is optionally substituted
with 1, 2, or 3 R₂₀₅ groups, wherein

20 each aryl or phenyl is optionally substituted with 1, 2,
or 3 groups that are independently R₂₀₅, R₂₁₀, or

C₁-C₆ alkyl substituted with 1, 2, or 3 groups that
are independently R₂₀₅ or R₂₁₀, and wherein

each heterocyclyl is optionally substituted with 1, 2, 3,
25 or 4 R₂₁₀;

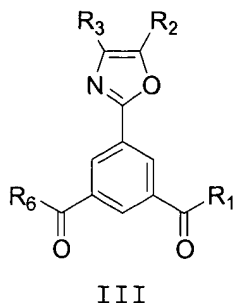
R₂₆₅ at each occurrence is independently -O-, -S- or -N(C₁-C₆
alkyl)-;

R₂₇₀ at each occurrence is independently R₂₀₅, halogen C₁-C₆
alkoxy, C₁-C₆ haloalkoxy, NR₂₃₅R₂₄₀, -OH, -C≡N, -CO-(C₁-C₄
30 alkyl), -SO₂-NR₂₃₅R₂₄₀, -CO-NR₂₃₅R₂₄₀, -SO₂-(C₁-C₄ alkyl), =O,
or

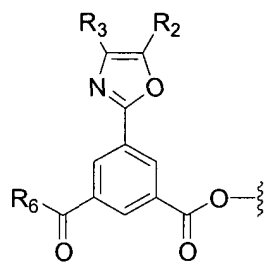
C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl or -(CH₂)₀₋₄-C₃-C₇
cycloalkyl, each of which is optionally substituted
with 1, 2, or 3 R₂₀₅ groups;

35 comprising

forming a reaction mixture comprising a compound of formula III

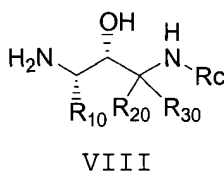


5 wherein



R₁ is OH, imidazolyl, halogen or
 R₂ and R₃ are independently H, phenyl, or C₁-C₄ alkyl; or
 R₂ and R₃ and the carbons to which they are attached form a
 benzene ring; and

10 R₆ is C₁-C₆ alkoxy or NR₄R₅; wherein
 R₄ and R₅ are independently C₁-C₆ alkyl;
 and a compound of formula VIII



15 in a solvent.

34. A process according to claim 33, wherein the solvent is selected from THF, DMF, CH₂Cl₂, and CHCl₃.

20 35. A process according to claim 35 wherein the reaction mixture comprises a base which is pyridine, collidine, di-tertiarybutyl pyridine, triethylamine, diisopropylethylamine, dimethylamino pyridine, or lutidine.

36. A process according to claim 35, wherein the reaction mixture further comprises an additive which is 1, 2, or 3 of the following:

EDCI, HOBT, benzotriazole, HOAT, HATU, or DCC.

5